























SWEDEN



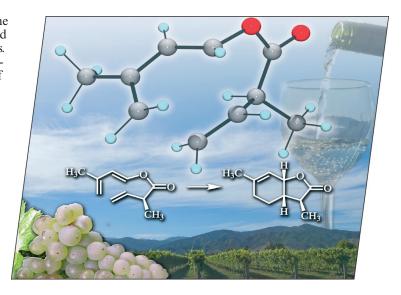




A union formed by chemical societies in Europe (ChemPubSoc Europe) has taken the significant step into the future by merging their traditional journals, to form two leading chemistry journals, the European Journal of Inorganic Chemistry and the European Journal of Organic Chemistry. Three further members of ChemPubSoc Europe (Austria, Czech Republic and Sweden) are Associates of the two journals.

COVER PICTURE

The cover picture shows a typical panoramic scene encountered in New Zealand: a vineyard situated near Blenheim at the base of the Southern Alps. Wine lactone, a sweet-smelling compound that contributes significantly to the aroma of a variety of white wines, is depicted in a reaction scheme in the foreground. The intramolecular Diels-Alder cycloaddition leading to wine lactone is diastereoselective, and the computationally predicted transition state is illustrated against the sky background. Scheurebe grapes and Riesling wine being poured into a glass are also depicted. Wine lactone is found in wines produced from both Scheurebe and Riesling grapes. Details are discussed in the article by M. A. Brimble et al. on p. 4405ff.



MICROREVIEW

Intramolecular Diels-Alder Reactions

K.-i. Tadano* 4381-4394

Natural Product Synthesis Featuring Intramolecular Diels-Alder Approaches - Total Syntheses of Tubelactomicins and Spiculoic Acid A

Keywords: Intramolecular Diels—Alder reaction / Natural products / Total synthesis / Cycloaddition / Tubelactomicins / Spiculoic acid A

- (+)-Tubelactomicin A: $R^1 = CO_2H$, $R^2 = Me$, $R^3 = Me$
- (+)-Tubelactomicin B: R¹ = Me, R² = Me, R³ = Me
- (+)-Tubelactomicin D: $R^1 = CO_2H$, $R^2 = CH_2OH$, $R^3 = Me$ (+)-Tubelactomicin E: $R^1 = CO_2H$, $R^2 = Me$, $R^3 = CH_2OH$
- o. Me gr

cins A, B, D, and E and the total synthesis of (+)-spiculoic acid A, all accomplished in the author's group by intramolecular Diels—Alder approaches, are summarized. The synthesis of (+)-tubelactomicin A by the Tatsuta group and the total synthesis of (-)-spiculoic acid A by the Baldwin/Lee group are also summarized.

The total syntheses of (+)-tubelactomi-

SHORT COMMUNICATIONS

Product-Selective Reaction

- A. Miyagawa, M. Naka, T. Yamazaki,* T. Kawasaki-Takasuka 4395–4399
- Unusual Behavior of the Anionic Species from (*E*)-1-Chloro-3,3,3-trifluoropropene (HCFC-1233t)

Keywords: Fluorine / Rearrangement / Alcohols / Alkenes

$$F_3C \xrightarrow{\hspace{1cm} \text{OH} \hspace{1cm} \text{i) MeLi/THF} \\ \hspace{1cm} \text{(<1.6 equiv)} \\ \hspace{1cm} \text{ii) PhCHO} \hspace{1cm} F_3C \xrightarrow{\hspace{1cm} \text{H} \hspace{1cm} \text{(>1.7 equiv)} \\ \hspace{1cm} \text{ii) PhCHO}} \hspace{1cm} F_3C \xrightarrow{\hspace{1cm} \text{OH} \hspace{1cm} \text{OH}} \hspace{1cm} CI \xrightarrow{\hspace{1cm} \text{ii) PhCHO}} \hspace{1cm} Ph$$

(*E*)-1-Chloro-3,3,3-trifluoropropene was smoothly deprotonated by MeLi at the position β to the CF₃ group, and exclusive formation of propargylic alcohols was observed by addition of appropriate carbonyl

compounds as long as up to 1.6 equiv. of MeLi was used, whereas more than 1.7 equiv. of the same base led to selective formation of allylic alcohols.

Hydrophosphinylation

- Regioselective Metal-Catalyzed Hydrophosphinylation of Alkynes: Synthesis of Enantiopure α or β -Substituted Vinylphosphane Oxides

Keywords: Alkynes / Homogeneous catalysis / Phosphorus heterocycles / Regioselectivity

Palladium was found to catalyze the regioselective Markovnikov addition of chiral enantiopure 1*r*-oxo-2*c*,5*t*-diphenylphospholane (1) to terminal alkynes, whereas

rhodium catalysis offers selectively the (E)-anti-Markovnikov adduct. This strategy offers rapid access to chiral and enantiopure α - or β -substituted-1-alkenylphospholanes.



FULL PAPERS

Wine lactone (1), a highly scented organic compound, has been synthesised by using an intramolecular Diels-Alder (IMDA) cycloaddition. Microwave irradiation of a range of ester-containing nonatrienes re-

sulted in a kinetically controlled distribution of isomeric bicyclo[4.3.0] lactones. Transition-state analysis by DFT at the B3LYP/6-31+G(d) level was in agreement with the experimental results.

Intramolecular Diels-Alder Reactions

P. D. O'Connor, U. B. Kim, M. A. Brimble* 4405–4411

Synthesis of (±) Wine Lactone and Its Analogues by a Diels-Alder Approach

Keywords: Diels—Alder reactions / Cycloaddition / Density functional calculations / Flavour / Lactones

Sonogashira Cross-Coupling

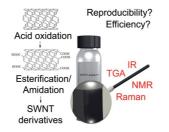
Without using any copper species and under robust conditions, rapid microwave-assisted Sonogashira cross-coupling of aryl iodides and bromides with terminal alkynes using Pd-EnCatTM TPP30 were conducted. Both electron-rich and electron-deficient aryl halides reacted smoothly with a broad variety of terminal alkynes in MeCN at $100-120\,^{\circ}\text{C}$.

Pd-EnCatTM TPP30 as a Catalyst for the Generation of Highly Functionalized Aryland Alkenyl-Substituted Acetylenes via Microwave-Assisted Sonogashira Type Reactions

Keywords: Cross-coupling / Heterogeneous catalysis / Reusable catalyst / Palladium / Microwave chemistry / Enynes

Carbon Nanotubes

Can we trust chemistry? Numerous methods for covalent functionalization of SWNTs have been reported in the literature. However, only a few studies have been concerned with the reproducibility and relative efficiency of these methods. Moreover, the fact that all batches of SWNTs and oxidized SWNTs are essentially different is sometimes neglected. This survey deals with these important topics.



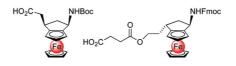
C.-H. Andersson, H. Grennberg* 4421–4428

Reproducibility and Efficiency of Carbon Nanotube End-Group Generation and Functionalization

Keywords: Nanotubes / Nanotechnology / Functionalization / Esterification / Amidation

Amino Acid Analogues

As a contribution to bioorganometallic chemistry, a dia- and enantioselective synthesis of novel carbocyclic amino acid analogues with a 1,2-ferrocenocyclopentene backbone has been developed.



A. Hunold, I. Neundorf, P. James, J. Neudörfl, H.-G. Schmalz* ... 4429–4440

Stereoselective Synthesis of New Ferrocene-Derived Amino Acid Building Blocks

Keywords: Metallocenes / Amino acids / Chirality / Stereochemistry / Solid-phase synthesis / Peptidomimetics

CONTENTS

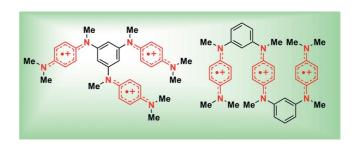
Polycationic Multispin Systems

A. Ito,* D. Sakamaki, H. Ino, A. Taniguchi, Y. Hirao, K. Tanaka,* K. Kanemoto,* T. Kato* 4441–4450



Polycationic States of Oligoanilines Based on Wurster's Blue

Keywords: Radical ions / Oxidation / Magnetic properties / Redox chemistry / EPR spectroscopy



The spin multiplicities of the dominant species of two oligoanilines based on Wurster's blue generated by three equivalents of oxidant have been found to exist as quartet and doublet states at low temperatures.

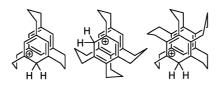
These results demonstrate that the intramolecular connectivity between the spincontaining units decisively influences the spin preference of the multispin systems based on oligoanilines.

Cationic Multibridged Cyclophanes



Stable-Ion NMR Spectroscopy and GIAO-DFT Study of Carbocations Derived from Multibridged [3_n]Cyclophanes

Keywords: Cyclophanes / Protonation / Carbocations / Donor—accepter systems / Superacidic systems



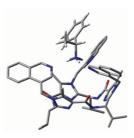
Stable-ion NMR spectroscopy and theoretical studies (GIAO-DFT and NICS) are reported for carbocations derived from multibridged [3n] (n = 3, 4, and 5) cyclophanes.

Enantiomeric Recognition

M. Schnopp, G. Haberhauer* ... 4458-4467

Highly Selective Recognition of α -Chiral Primary Organoammonium Ions by C_3 -Symmetric Peptide Receptors

Keywords: Peptides / Receptors / Macrocyclic ligands / Enantioselectivity / Organoammonium ions

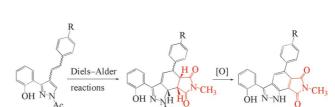


Chiral macrocyclic imidazole peptides with different binding arms were synthesized. The investigation of their ability to bind α -chiral primary organoammonium ions in CDCl₃ shows that these cyclopeptides can be used as excellent and highly selective receptors for enantiomeric discrimination.

Diels-Alder Reactions

Synthesis of New 1*H*-Indazoles through Diels-Alder Transformations of 4-Styryl-pyrazoles under Microwave Irradiation Conditions

Keywords: Diels-Alder reaction / Microwave irradiation / Indazoles / Cycloaddition / Nitrogen heterocycles / Dehydrogenation



Diels—Alder cycloaddition reactions between (E)- or (Z)-1-acetyl-3-(2-hydroxyphenyl)-4-styrylpyrazoles and N-substituted maleimides under microwave irradia-

E and Z isomers

tion and solvent-free conditions yield tetrahydroindazoles; the latter can be dehydrogenated to give the corresponding indazole-type compounds.



Unsymmetric Tweezer Receptors

The synthesis of an orthogonally protected template 1 for the solid-phase synthesis of unsymmetrical tweezers is presented. The synthesis consists of seven steps with an

overall yield of 34%. The key step is the selective mono reduction of a symmetric diazide taking advantage of a biphasic reaction mixture.

H. Y. Kuchelmeister, C. Schmuck* 4480–4485

An Efficient Synthesis of an Orthogonally Protected Aromatic Diamine as Scaffold for Tweezer Receptors with Two Different Arms

Keywords: Protecting groups / Template synthesis / Peptides / Solid-phase synthesis / Receptors

Immobilized Organocatalysts

A new strategy for the immobilization of iminium organocatalysts has been developed through the acid-base assembly of

multidentate chiral primary amines and solid polyacids and has been used in the Diels-Alder reactions of α -substituted acroleins.

J. Li, X. Li, P. Zhou, L. Zhang, S. Luo,* J.-P Cheng* 4486-4493

Chiral Primary Amine—Polyoxometalate Acid Hybrids as Asymmetric Recoverable Iminium-Based Catalysts

Keywords: Polyoxometalates / Amines / Asymmetric catalysis / Supported catalysts / Diels—Alder reactions

The formation of resin-bound precursors of natural products and the subsequent cleavage of these substances are described. The common structural motif of the synthesized natural products is a diaryl ether functionality that was built up via an Ullmann-type reaction of phenols and aryl halides on solid phases.

SPOS of Natural Products

N. Jung, S. Bräse* 4494-4502

Synthesis of Natural Products on Solid Phases via Copper-Mediated Coupling: Synthesis of the Aristogin Family, Spiraformin A, and Hernandial

Keywords: Solid-phase synthesis / Combinatorial chemistry / Ullmann coupling / Natural products / Synthetic methods / Aristogins / Diaryl ether

Green Ways to Indolylchromans

An efficient green protocol for the synthesis of indolyl(nitro)chromen derivatives by employing "on-water" conditions is re-

ported. The factors influencing the diastereoselective addition of various indoles with nitrochromenes have been studied.

"On-Water"-Promoted *C*-Alkylation of Indoles with 2-Aryl-3-nitro-2*H*-chromenes under Catalyst-Free Conditions

Keywords: "On-water" chemistry / Alkylation / 2*H*-Chromenes / Indoles / Green chemistry / Synthetic methods

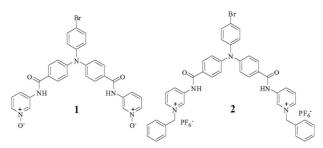
CONTENTS

Anion Recognition

K. Ghosh,* G. Masanta, A. P. Chattopadhyay 4515-4524

Triphenylamine-Based Pyridine N-Oxide and Pyridinium Salts for Size-Selective Recognition of Dicarboxylates

Keywords: Receptors / Anions / Anion recognition / Fluorimetric detection



Triphenylamine-based receptors 1-2 have been designed and synthesized for size-selective recognition of dicarboxylates. Binding takes place at charged sites with concomitant PET-based (photo-induced

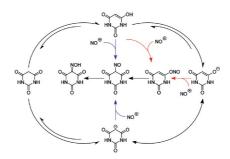
electron transfer) quenching of emission of triphenylamine motif. Receptor 2 shows more effective binding than 1 and is selective for pimelate.

Ambident Nucleophiles



Enol Nitrosation Revisited: Determining Reactivity of Ambident Nucleophiles

Keywords: Enols / Ketones / Nitrosation / Reaction mechanisms / Tautomerism



An alternative methodology for determining chemical reactivity of ambident nucleophiles is reported. This approach is based on the different operating mechanisms for enol nitrosation.

Switchable Fluorescent Chemosensor



On-Off Switchable Binuclear Chemosensor Based on Thiacalix[4]crown Armed with Pyrene Moieties

Keywords: Thiacalixarenes / Crown compounds / Pyrene / Fluorescence / Molecular devices / Chemosensor

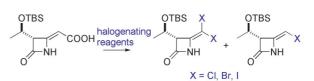


A new fluorescent "on-off" switchable chemosensor based on a thiacalix[4]arene with a 1,3-alternate conformation and two different types of cation binding sites has been synthesized.

Halodecarboxylation of β-Lactams

Halodecarboxylation Reaction of 4-Alkylidene- β -lactams

Keywords: Nitrogen heterocycles / Lactams / Halodecarboxylation / Halogenation / Alkenes



A Hunsdiecker-type halodecarboxylation reaction has been used for the synthesis of a new family of 4-alkylidene- β -lactams. The scope and limits of the synthesis of

(chloro-, bromo-, and iodoalkylidene)azet-idinones are explored. The unexpected formation of dihalovinyl derivatives was investigated by ¹H NMR analysis.

Supporting information on the WWW (see article for access details).

If not otherwise indicated in the article, papers in issue 25 were published online on August 11, 2009

^{*} Author to whom correspondence should be addressed.